- **62**. A composition in pharmaceutical dosage form comprising an anticancer effective amount comprising a compound or pharmaceutically acceptable salt of claim **48**, in combination with a pharmaceutically acceptable carrier, additive, or excipient.
- **63**. The composition of claim **62**, comprising a compound of the chemical structure

$$\begin{array}{c} \text{HO} \\ \text{TFA} \\ \text{HO} \\ \text{HO}$$

or a pharmaceutically acceptable salt thereof.

- **64.** The composition according to claim **62**, wherein said UPA is human recombinant urokinase.
- **65**. The composition according to claim **62** further comprising an additional anticancer agent.
- **66.** The composition according to claim **65** wherein said additional anticancer agent is an antimetabolite, an inhibitor of topoisomerase I and II, an alkylating agent, a microtubule inhibitor or a mixture thereof.
- **67**. The composition according to claim **65** wherein said additional anticancer agent is a FLT-3 inhibitor, a VEGFR inhibitor, an EGFR TK inhibitor, an aurora kinase inhibitor,

a PIK-1 modulator, a Bcl-2 inhibitor, an HDAC inhibitor, a c-MET inhibitor, a PARP inhibitor, a Cdk inhibitor, an EGFR TK inhibitor, an IGFR-TK inhibitor, an anti-HGF antibody, a PI3 kinase inhibitors, an AKT inhibitor, a JAK/STAT inhibitor, a checkpoint-1 or 2 inhibitor, a focal adhesion kinase inhibitor, a Map kinase (mek) inhibitor, a VEGF trap antibody or a mixture of any of the foregoing agents.

68. A method of treating cancer in a patient in need comprising administered to said patient an effective amount of a composition according to claim **48**.

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